Established Condition & Commonly Used ICD-10 Codes v. 7/17/17

Est Cond?	ICD-10	Diagnosis for Birth Mandate / Dually Eligible for Part C & B
Yes	H93.3X9	Auditory Neuropathy Spectrum Disorder (ANSD) (Disorder of Acoustic Nerve)
Yes	F84.0	Autism Spectrum disorder
Yes	H54	Blindness NOS
Yes	H90.2	Conductive hearing loss, unspecified
Yes	H47.619	Cortical Blindness, unspecified
Yes	H90.8	Mixed conductive and sensorineural hearing loss, unspecified
Yes	H90.5	Sensorineural Hearing Loss (Unspec)
Yes	H90.4	Sensorineural hearing loss, unilateral
Yes	H91.93	Unspecified hearing loss - Bilateral

Note: "*" = follow until 12 months post referral

Changes post 6/30/17 meeting

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Est Cond?	ICD-10	Diagnosis for Part C Established Condition
No	R29.2	Abnormal Reflex
No	Q10.4	Abs & agenesis of lacrimal apparatus
No	Q13.1	Absense of IRIS
No	Q77.4	Achondroplastic Dwarfism
Yes	Q87.0	Acrocephalosyndactyly type III (aka Saethre–Chotzen syndrome)
No	J05.0	acute obstructive laryngitis
No	H65.00	Acute serous otitis media, unspec ear
No	I62.01	Acute Subdural Hemorrhage
No	I50.43	Acute Systolic and Diastolic Cong. Heart Failure
No	Z90.49	Acquired Absensce of Other Specified Part of Digestive Tract
No	P02.5	Affected by Compression of Umbilical Cord (NEC)
No	P02.5	Affected by Entanglement (knot) of Umbilical Cord (NB)
No	P02.5	Affected by Umbilical Cord (Tightly) (NB)
Yes	Q04.0	Agenesis of Corpus Callosum*
No	E70.30	Albinism, unspecified
No	T 78.40	Allergy Unspecified
Yes	S68.412S	Amputation of Left Hand @ Wrist
Yes	S68.411S	Amputation of Right Hand @ Wrist
No	D55.0	Anemia due to glucose - 6 - phosphate dehydrogenase (G6PD) deficiency
No	P61.2	Anemia of prematurity
No	R63.0	Anorexia
Yes	G93.1	Anoxic brain damage
No	R48.5	Apraxia
No	G93.0	Arachnoid Cyst
Yes	Q07.02	Arnold-Chiari syndrome with hydrocephalus
Yes	Q74.3	Arthrogryposis

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Est Cond?	ICD-10	Diagnosis for Part C Established Condition
No	Q28.2	Arteriovenous malform. of cereb. vessels
No	R53.1	Asthenia NOS
No	R27.0	Ataxia - Unspecified
No	Q21.1	Atrial Septal defect
No	Q21.2	Atrioventricular septal defect
Yes	G60.0	Atrophy, Charcot-Marie-tooth Syndrome
No	P36.9	Bacterial Sepsis of Newborn
Yes	E75.4	Batten-Mayou Disease
No	E53.8	B-Complex Deficiency
No	F91.8	Behavior - Aggressive Behavior Disorder
No	F91.0	Behavior – Conduct Disorder Confined to Family Context
No	F94.1	Behavior – Reactive Attachment Disorder of Childhood
No	F91.1	Behavior - Solitary Aggressive Disorder
No	F91.8	Behavior - Tantrum-Conduct/Behavior Disorder
No	F91.1	Behavior - Un-socialized Aggressive Disorder
No	F91.1	Behavior – Un-socialized Conduct Disorder
No	K40.20	Bilat ing hernia, w/o obstr, NEC
No	P84	Birth Asphyxia
No	P15.9	Birth Trauma
Yes	Z94.81	Bone Marrow Transplant*
Yes	A48.51	Botulism (Infant)*
No	R00.1	Bradycardia, unspecified
No	Q04.6	Brain Anomaly (Cysts/Congenital)
No	G93.9	Brain Condition - NOS
Yes	P27.1	Bronchopulmonary dysplasia (BPD)* - Only if Oxygen dependent
No	R01.1	Cardiac Murmur, unspecified
Yes	I42.8	Cardiomyopathy (Primary)*
Yes	I63.50	Cerebral Artery Occlusion NOS
Yes	P10.1	Cerebral Hem. due to birth injury* (grade IV bleed)
Yes	G80.9	Cerebral Palsy, unspecified
Yes	G80.0	Cerebral Spastic Quadriplegia
Yes	E75.4	Ceroid-Lipofuscinosis, Neuronal
No	Q30.0	Choanal Atresia
No	Q99.9	Chromosomal abnormality, unspec
Yes	Q97.0	Chromosomal Anomalies – Karyotype 47, XXX
Yes	J44.9	Chronic/Obstructive Lung Disease
No	D66	Classic Hemophilia, NOS, A, Defic Factor VIII
Yes	E70.0	Classical phenylketonuria
Yes	Q36.8	Cleft Lip (Unilateral)* post repair
Yes	Q37.9	Cleft Lip and Palate* post repair
Yes	Q35.9	Cleft Palate* post repair
Yes	Q25.1	Coarctation of Aorta (pre/postductal)* only if surgically repaired
No	P04.41	Cocaine Exposed (prenatally)

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Est Cond?	ICD-10	Diagnosis for Part C Established Condition
No	H47.319	Coloboma of Optic Disc
No	Q20.0	Common arterial trunk
Yes	H90.2	Cond. hearing loss, unspecified
Yes	G80.8	Cong Cerebral Diplegia / Hemiplegia
Yes	P35.1	Cong Cytomegalovirus Infection (only if symptomatic)
No	Q65.1	Cong dislocation of hip, bilateral
No	Q65.0	Cong dislocation of hip, unilateral
Yes	Q75.9	Cong Malform of Skull and Face Unspec*
Yes	Q24.9	Cong Malform of the Heart Unspec* only if surgery required
No	Q89.9	Cong. Anomaly/deformity (NOS)
No	Q89.9	Cong. Anomaly/Lymphatic Malform (NOS)
No	Q15.9	Cong. Anomaly/Malform of Eye Unspec.
No	Q14.2	Cong. malformation of optic disc
No	Q39.2	Cong. tracheo-esoph fistula w/out atresia
No	Q87.1	Congenital Anomaly, Other specified
No	E34.3	Congenital Dwarfism
Yes	Q07.9	Congenital Encephalopathy*
No	P83.5	Congenital Hydrocele
No	Q62.0	Congenital hydronephrosis
Yes	Q07.9	Congenital Hypomyelination affecting Meningeal bands or folds
yes	E03.1	Congenital Hypothyroidism*
No	Q20.9	Cong. Malf. of cardiac chambers and connections, unspecified
Yes	Q16.4	Congenital Malformation of middle ear NOS
No	Q14.1	Congenital malformation of retina
Yes	Q17.9	Congenital Malformations of Ear NOS
No	D74.0	Congenital methemoglobinemia
Yes	Q23.3	Congenital Mitral Insufficiency* (only if severe)
Yes	H55.01	Congenital nystagmus
No	M41	Congenital Scoliosis
No	H50.60	Congenital Strabismus
No	Q32.0	Congenital tracheomalacia
No	P25.1	Congenital/Newborn Pneumothorax
Yes	R56.9	Convulsion NOS* only with ongoing medication treatment
No	R56.9	Convulsions (Idiopathic)
Yes	P90	Convulsions of Newborn*
Yes	Q20.8	Cor Biloculare (Cong. Malf. of Cardiac Chambers and Connections)
Yes	H47.619	Cortical Blindness, unspecified
No	P28.2	Cyanotic attacks of Newborn
Yes	E84.9	Cystic Fibrosis, unspecified
No	H55.89	Deficiency / Smooth pursuit movements (eye)
No	Q66.9	Deformity, foot, congenital NOS
No	E86.0	Dehydration
No	R62.0	Delay milestones (Late talker/walker) – NEC

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Est Cond?	ICD-10	Diagnosis for Part C Established Condition
Yes	Z99.11	Dependent on Respirator or Ventilator
No	L27.9	Dermatitis due to Ingested substance
No	L27.9	Dermatitis due to unspec. substance taken internally
No	F82	Developmental Dyspraxia
Yes	E23.2	Diabetes Insipidus
Yes	E11.9	Diabetes, Diabetic - unspecified
No	R19.7	Diarrhea, unspecified
No	Q20.5	Discordant atrioventricular connection
No	Q20.3	Discordant ventriculoarterial connection
No	M24.20	Disorder of ligament, unspecified site
No	M62.9	Disorder of Muscle, unspecified
Yes	Q20.4	Double inlet ventricle
Yes	Q20.2	Double Outlet left ventricle
Yes	Q20.1	Double outlet right ventricle
Yes	Q90.9	Down Syndrome
No	R13.10	Dysphagia Unspecified
No	J15.7	Eaton's Agent (Pneumonia)
Yes	Z92.81	ECMO - Extracorporeal Membrane Oxygenation
No	Q82.4	Ectodermal dysplasia (anhidrotic)(Cong)
Yes	G04.90	Encephalitis (postnatally acquired)*
Yes	Q01	Encephalocele
Yes	G93.40	Encephalopathies (Degenerative) NOS
No	Q85.8	Encephalotrigeminal Angiomatosis
Yes	G40.823	Epil spasms, intract, w/ status epilepticus
Yes	G40.824	Epil spasms, intract, w/o status epilepticus
Yes	G40.919	Epilepsy, unspec, intractable, w/o status epilepticus
Yes	G40.911	Epilepsy, unspec, intractable, w/status epilepticus
No	P14.0	Erb's Paralysis due to birth injury
No	P07.14	Ex low weight nb, less 1000-1249 gr
No	P07.15	Ex low weight nb, less 1250-1499 gr
Yes	P07.01	Ex low weight nb, less 500 gr*
Yes	P07.02	Ex low weight nb, less 500-749 gr*
Yes	P07.03	Ex low weight nb, less 750-999 gr*
No	Q79.2	Exomphalos
Yes	Q45.8	Exstrophy, Abdominal Contents*
Yes	Q64.10	Exstrophy, Bladder*
No	G25.9	Extrapyramidal & movement disorder, unspec
No	P11.3	Facial Nerve Paralysis (Congenital)
No	R62.51	Failure to thrive (Child)
No	P92.6	Failure to thrive in newborn
No	E43	Failure to Thrive. Unspec severe protein calorie maln.
No	R63.4	Failure to Thrive: Ab. loss of weight
No	R63.3	Feeding difficulties for infant / child

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Est Cond?	ICD-10	Diagnosis for Part C Established Condition
Yes	Q86.0	Fetal Alcohol Syn. (dysmorphic)
Yes	Q99.2	Fragile X Syndrome
Yes	Q75.4	Franceschetti–Klein (Wildervanck) Syndrome
Yes	E74.21	Galactosemia*
No	K21.0	Gastro-esoph reflux dis w esophagitis
No	K21.9	Gastro-esoph reflux dis w/o esophagitis
Yes	Z93.1	Gastronomy tube (G-Tube) Dependent
No	Q79.3	Gastroschisis
Yes	E75.22	Gaucher Disease (Lipidoses)
No	H42	Glaucoma in diseases (classified elsewhere)
Yes	E74.00	Glycogen storage disorder
No	I50.9	Heart failure, unspecified
Yes	Z94.1	Heart Transplant Status
No	D18.0	Hemangioma NOS
Yes	D18.02	Hemangioma of intracranial struct.
No	D18.01	Hemangioma of skin and subqu
No	D18.1	Hemolymphangioma
No	P12.2	Hemorrhage/Subgaleal (Epicranial Subaponeurotic Hem. Due to Birth Injury)
No	I78.0	Hereditary hemorrhagic telangiectasia
Yes	P91.660	HIE/Body Cooling (Hypoxic Ischemic Enceph.)*
No	Q43.1	Hirschsprung's Disease
Yes	Q04.2	Holoprosencephaly
No	B20	Human Immunodeficiency Virus (HIV)
Yes	E76.02	Hurler-Scheie Syndrome
Yes	G91.9	Hydrocephalus (shunt)*
Yes	G91.1	Hydrocephalus, Obstructive
Yes	Q03.8	Hydrocephalus, Other Congenital
No	D18.1	Hygroma (congenital) (cystic)
No	E83.52	Hypercalcemia
No	R20.3	Hyperesthesia of Skin
No	H52.03	Hypermetropia, bilateral
Yes	I11.0	Hypertensive Heart Disease w/ Heart Failure
No	P94.1	Hypertonicity
No	J35.1	Hypertrophy of tonsils
Yes	R20.1	Hypoesthesia of Skin
Yes	E16.2	Hypoglycemia - Unspecified (only if severe or chronic)
Yes	Q23.4	Hypoplastic Left Heart Syndrome
Yes	Q22.6	Hypoplastic Right Heart Syndrome
No	E87.1	Hyposmolality and hyponatremia
No	Q54.9	Hypospadias, unspecified
No	E03.9	Hypothyroidism, unspec
No	D84.9	Immunodeficiency, Unspec
No	Q42.3	Imperforate Anus (Cong. absence of anus)

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Est Cond?	ICD-10	Diagnosis for Part C Established Condition
No	K40.90	Inguinal Hernia
No	H55.89	Irregular Eye Movements
Yes	I61.9	Intracerebral (nontraumatic) Hem.*
Yes	G06.0	Intracranial abscess and granuloma
No	P52.3	Intravent. Hem. in prem infant (non-traum)
Yes	P52.0	Intraventricular Hem./Newborn, only Grade III & IV
No	Q20.6	Isomerism of atrial appendages
Yes	E75.4	Jansky-Bielschowski Amaurotic Idiocy
Yes	E75.23	Krabbe disease
Yes	E75.4	Kuf's Disease
Yes	C96.6	LCH Langerhans Cell Histiocytosis
No	Q31.5	Laryngomalacia (Congenital)
Yes	Z77.011	Lead Exposure (Toxic) If level 10 or higher*
Yes	Q23.3	Leaking Heart Valve – Cong. Mitral Insufficiency, if severe
Yes	G04.81	Leukoencephalitis
No	D18.1	Lymphangioendothelioma
No	D18.1	Lymphangioma (Cap., Cavernous, Cystic)
No	D18.1	Lymphangiomyoma
No	Q38.2	Macroglossia, Congenital
No	K90.9	Malabsorption NEC
Yes	Q04.9	Malformation of Brain (Cong/Unspec)*
No	Q24.5	Malformation of coronary vessels
Yes	C71.8	Malig neoplasm of overlapping sites of brain
No	Q75.4	Mandibulofacial Dysostosis
Yes	E71.0	Maple-Syrup-Urine Disease
Yes	Q87.40	Marfan Syndrome
No	E84.11	Meconium ileus in cystic fibrosis
No	P76.0	Meconium Plug Syndrome (Nb)
Yes	Q04.5	Megalencephaly
Yes	G00.2	Meningitis – Streptococcal*
Yes	A02.21	Meningitis due /Salmonella infect.*
Yes	G03.9	Meningitis NOS*
No	E87.4	Metabolic Acidosis - W/ respiratory acidosis
No	E87.2	Metabolic Acidosis NEC
No	P74.0	Metabolic Acidosis of Newborn (Late)
Yes	Q02	Microcephalus, microcephaly (under 3%)
Yes	Q04.3	Microgyria
Yes	Q17.2	Microtia
No	E44.1	Mild Protein - calorie Malnutrition
Yes	H90.8	Mixed cond & sensorineural hearing loss, unspec
No	F98.4	Movement disorder (Head Banging)
No	F98.4	Movement disorder (rocking)
No	A49.02	MRSA - Methicillin resistant Staphylococcus aureus inf.

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Est Cond?	ICD-10	Diagnosis for Part C Established Condition
No	A49.01	MSSA - Methicillin susceptible Staphylococcus aureus inf.
No	Q89.7	Multiple congenital malformations, not elsewhere classified
No	R53.1	Muscle Weakness NOS
Yes	G71.0	Muscular Dystrophy
Yes	I21.3	Myocardial Infarction Unspec (STEMI)
Yes	G70.9	Myoneural disorder
Yes	G71.11	Myotonic Dystrophy
Yes	P96.1	NAS - Neonatal Abstinence Syndrome* (only if permanent effects)
Yes	Q01.1	Nasofrontal Encephalocele*
No	P05.15	NB small for gest age 1250-1499 gr
No	P05.14	NB small for gest age, 1000-1249 gr
No	P05.16	NB small for gest age, 1500-1749 gr
No	P05.17	NB small for gest age, 1750-1999 gr
Yes	P05.12	NB small for gest age, 500-749 gr*
Yes	P05.13	NB small for gest age, 750-999 gr*
Yes	P05.11	NB small for gest age, less 500 gr*
No	P05.10	NB small for gest age, unspec wt
Yes	P77.9	Necrotizing enterocolitis/NB, unspec* (only if post surgical)
No	P59.0	Neo jaundice assoc.with preterm del
No	P24.9	Neonatal aspiration, unspecified
No	P91.4	Neonatal cerebral depression
No	P91.0	Neonatal cerebral ischemia
Yes	P91.2	Neonatal cerebral leukomalacia
No	P24.81	Neontal asp. with pneumonitis NEC
No	P01.5	Newborn affected by mult preg
No	P28.4	Newborn NEC Apnea - Spells of
No	I60.9	Non-trauma subarach hem., unspec
Yes	G91.1	Obstructive Hydrocephalus
No	Q79.2	Omphalocele
Yes	H47.039	Optic Nerve Hypoplasia Unspec
Yes	Q78.0	Osteogenesis imperfecta
No	E88.9	Other and unspec metabolic disorders
No	L20.89	Other Atopic Dermatitis
No	Q23.8	Other cong malform of aortic & mitral valves
No	Q44.7	Other cong malformations of liver
No	P61.4	Other cong. anemias, NEC
Yes	Q03.8	Other Congenital Hydrocephalus
Yes	Q79.59	Other congenital malformations of abdom wall*
Yes	Q93.5	Other del/part of chromo, appl to Angelman Synd
No	E23.6	Other disorders of pituitary gland
No	R20.8	Other Disturbances of Skin Sensations
No	F98.29	Other feeding disorders of infancy
No	P92.8	Other feeding problems of newborn

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Est Cond?	ICD-10	Diagnosis for Part C Established Condition
No	D55.8	Other Hemolytic anemias due to enzyme defic
No	D69.49	Other primary thrombocytopenia
No	Q87.89	Other spec cong malform synd, NEC
No	S06.890A	Other spec intracranial injury w/o loss of consc, initial enc
No	Q65.89	Other spec. cong. deformities of hip
Yes	Q04.8	Other Specified Congenital Malformation of the Brain*
No	F91.8	Other specified Disruptive, impulse control, and conduct disorder
No	A41.89	Other specified sepsis
No	D56.8	Other thalassemias
Yes	Z99.81	Oxygen Dependent (Supplemental)
No	Q25.0	Patent Ductus Arteriosus
No	P29.3	Persistent Fetal Circulation
Yes	P04.1	Phenytoin Exposed (Prenatally)
No	F98.3	Pica of infancy and childhood
Yes	Q87.0	Pierre Robin Synd - Cong. Malform synd predom aff face
No	J15.7	Pleuro-Pneumonia-Like-Organism (PPLO)
No	J15.7	Pneumonia Microplasm
No	J15.7	Pneumonia/ Acute
Yes	Q04.3	Polymicrogyria
Yes	Q04.6	Porencephalic Cysts (Congenital)
No	Q18.1	Preauricular Cyst/congenital/fistula
No	P07.21	Preterm < 23 Comp (23 wks 0 days)
No	P07.22	Preterm 23 Comp (23wks 0d thru 23wks 6d)
No	P07.23	Preterm 24 Comp (24wks 0d thru 24wks 6d)
No	P07.24	Preterm 25 Comp (25wks 0d thru 25wks 6d)
No	P07.25	Preterm 26 Comp (26wks 0d thru 26wks 6d)
No	P07.26	Preterm 27 Comp (27wks 0d thru 27wks 6d)
No	P07.31	Preterm 28 Comp (28wks 0d thru 28wks 6d)
No	P07.32	Preterm 29 Comp (29wks 0d thru 29wks 6d)
No	P07.33	Preterm 30 Comp (30wks 0d thru 30wks 6d)
No	P07.34	Preterm 31 Comp (31wks 0d thru 31wks 6d)
No	P07.35	Preterm 32 Comp (32wks 0d thru 32wks 6d)
No	P07.36	Preterm 33 Comp (33wks 0d thru 33wks 6d)
No	P07.37	Preterm 34 Comp (34wks 0d thru 34wks 6d)
No	P07.38	Preterm 35 Comp (35wks 0d thru 35wks 6d)
No	P07.39	Preterm 36 Comp (36wks 0d thru 36wks 6d)
No	I27.0	Primary Pulmonary Hypertension
No	F50.8	Psychogenic Vomiting
Yes	Q22	Pulmonary Valve Atresia
No	Q04.3	Reduction Deformities of the Brain
No	P92.1	Regurgitation of newborn
Yes	Q60.2	Renal agenesis, bilateral, unspecified
No	R09.2	Respiratory arrest

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Est Cond?	ICD-10	Diagnosis for Part C Established Condition
No	P22.0	Respiratory distress syndrome of nb
No	J96.90	Respiratory failure, unspecified whether with hypoxia or hypercapnia
Yes	H35.109	Retinopathy of Prematurity, unspec elig., (Grade 3 and above only)
Yes	P35.0	Rubella (Prenatally Acquired)
Yes	Q87.0	Saethre–Chotzen syndrome (aka Acrocephalosyndactyly type III)
Yes	Q04.6	Schizencephaly
No	L21.1	Seborrheic infantile dermatitis
Yes	R56.9	Seizures NOS, only if ongoing medication / treatment
No	A41.9	Sepsis - Unspecified
No	A02.1	Sepsis due to Salmonella
No	A41.9	Septicemia - Unspecified
Yes	T74.4	Shaken Infant Syndrome
No	D57.1	Sickle Cell disease without crisis
No	D57.3	Sickle Cell trait
No	P28.3	Sleep (Obst) (Primary) - Apnea
Yes	Q05.4	Spina Bifida
Yes	G12.0	Spinal Muscular Atrophy (Werdnig-Hoffman)
No	I21.19	ST elevation (STEMI) myocardial infarction
No	Q25.6	Stenosis of Pulmonary Artery
Yes	A53.9	Syphilis (Prenatally acquired)
No	Q66.6	Talipes Valgus (Congenital)
No	Q66.3	Talipes Varus (Congenital
Yes	Q87.2	TAR Syndrome: (Thrombocytopenia with Absent Radius Syndrome)
Yes	E75.02	Tay Sachs (Nervous System Disorder)
Yes	Q21.3	Tetralogy of Fallot*
No	D69.42	Thrombocytopenia/ Cong/hereditary
No	F93.8	Timidity, Child
No	M43.6	Torticollis
Yes	B58.9	Toxoplasmosis
No	Q31.5	Tracheo/Laryngomalacia
No	Q39.1	Tracheo-esoph fistula w/ atresia of Esophagus
Yes	Z93.0	Tracheostomy Status
Yes	S06.2X0D	Traumatic Brain Injury, Diffused w/o LOC
Yes	Q75.4	Treacher Collins Syndrome
No	R25.1	Tremor unspecified
Yes	Q97.0	Triple X Syndrome (Chromosomal Anom)
Yes	Q91.7	Trisomy 13 (Patau's Syndrome) Unspec
Yes	Q91.3	Trisomy 18 (Edwards Syndrome) Unspec
Yes	Q85.1	Tuberous sclerosis*
Yes	Q96.9	Turner-Ullrich Syndrome
No	K42.9	Umbilical Hernia
No	R20.9	Unspec .Disturbances of Skin Sensation
No	T 74.92	Unspec Confirmed Child Maltreat

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Est Cond?	ICD-10	Diagnosis for Part C Established Condition
No	H93.099	Unspec degen and vascular dis of unspec ear
No	H72.90	Unspec perf. of tympanic membrane
No	H02.409	Unspec. ptosis of unspec eyelid
No	H50.00	Unspecified esotropia
No	H50.50	Unspecified Heterophoria
No	N19	Unspecified kidney failure
No	R27.9	Unspecified lack of coordination
No	H55.00	Unspecified nystagmus
Yes	P39.9	Varicella (Perinatally acquired)
No	Q21.0	Ventricular septal defect
No	H54.7	Vision Loss Unspecified
Yes	A92.5	Zika Disease only with Microcephaly* (not just exposure)

Est Cond?	ICD-10	Common Diagnosis for Developmental Delay - Not Estab Condition
No	R48.2	Apraxia
No	F80.89	Artic delay - Other Dev Disorder of Speech & Language
No	F80.0	Artic delay - Phonological Disorder
No	H93.25	Central auditory proc disorder/ Rec. Delay
No	F88	Cognitive Developmental Delay
No	F88	Cognitive Developmental Delay (Borderline)
No	R62.0	Delayed milestones (Late talker/Late Walker) – NEC
No	F82	Dev Delay for Motor Coordination
No	F80.1	Expressive Language Disorder
No	F80.1	Expressive type dev dysphasia
No	F88	Global Developmental Delay
No	F82	Gross and/or Fine motor delay
No	R25.9	Involuntary movement disorder
No	F88	Mixed Developmental delay skills
No	F80.2	Mixed rec-exp lang disorder
No	R49.8	Other voice & resonance disorders
No	R47.81	Slurred Speech/ Speech disturb.
No	F80.81	Stuttering Childhood onset
No	F80.4	Speech Delay due to Hearing Loss
No	F80.9	Speech Delay Unspec
No	R27.9	Unspecified lack of coordination (Gross motor and/or Fine motor)